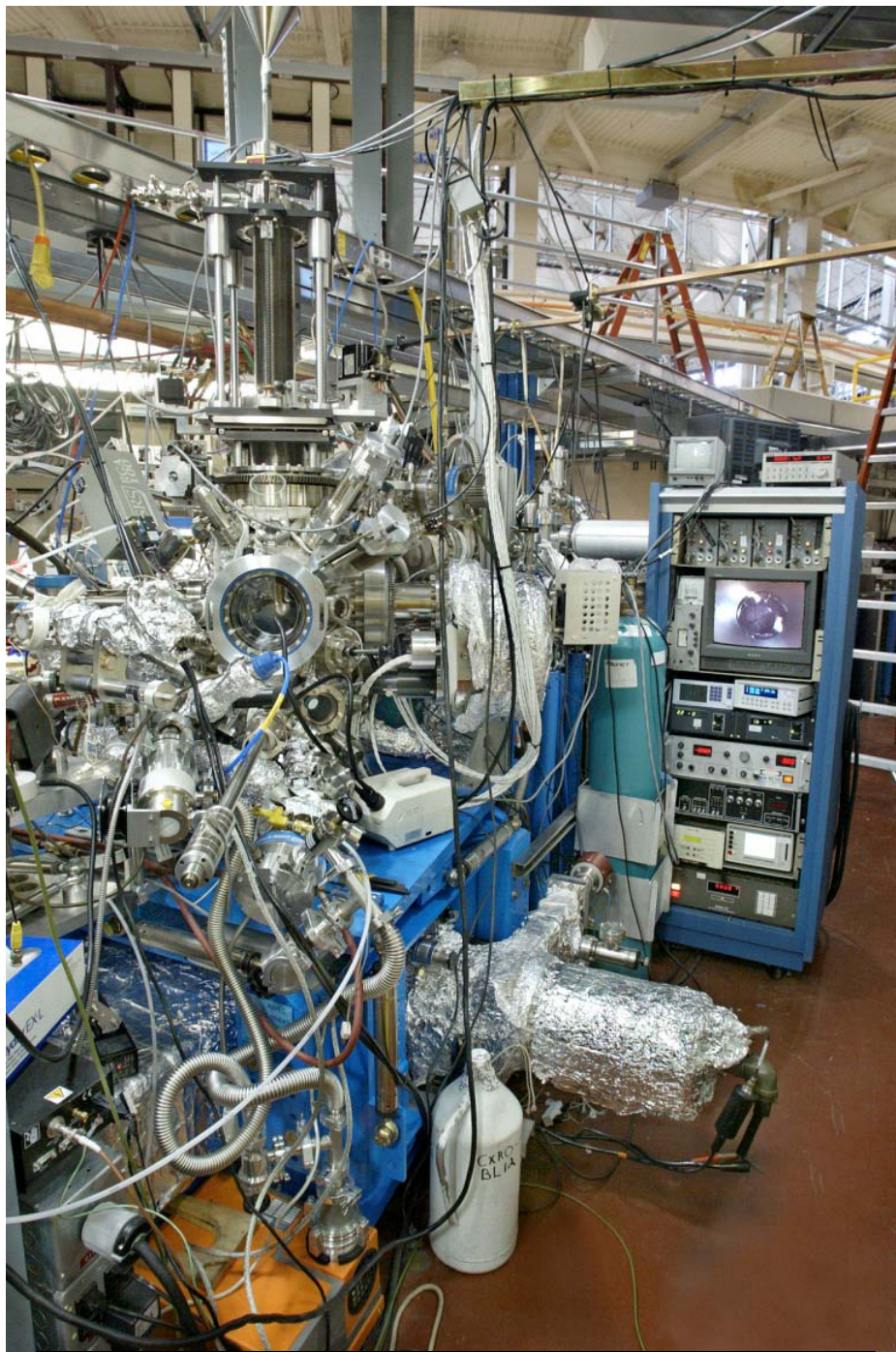


# Beamline 12.0.1.1 Manual

(Note: Always check end of this document for recent appendices)



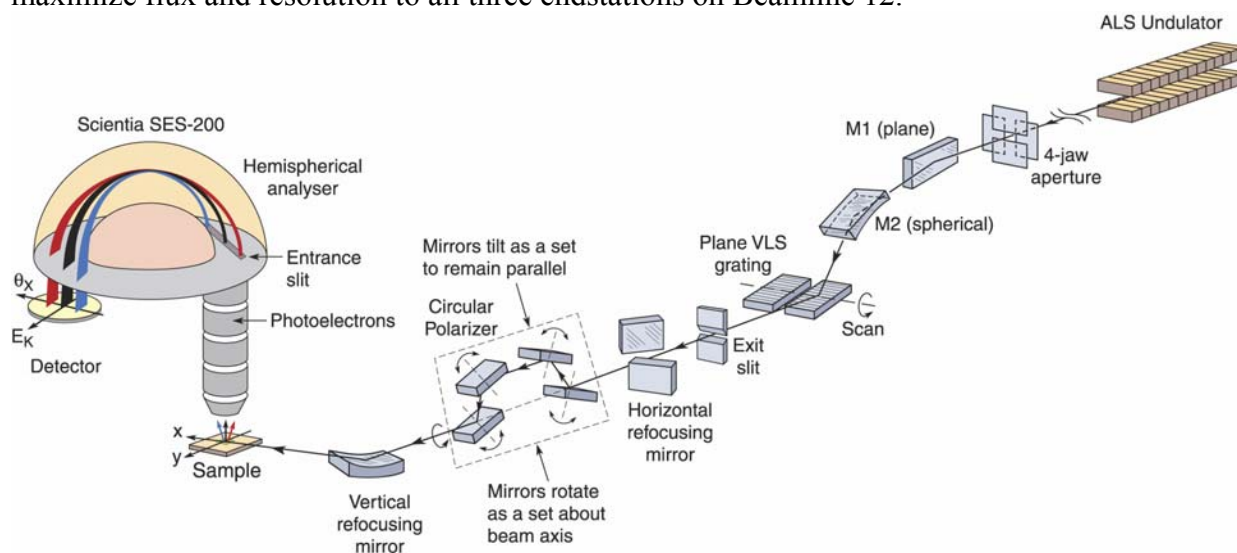
Overview of main experimental chamber at Beamline  
12.0.1.1

## ***Table of Contents:***

- Overview of the Beamline...  
**Page 2**
- Photoemission Spectroscopy...  
**Page 3**
- The Circular Polarizer...  
**Page 4**
- Vacuum Safety Interlock System  
**Page 6**
- Beam Detection  
**Page 7**
- Swapping the Sample ...  
**Page 8**
- Venting the Main Chamber...  
**Page 11**
- Using the LEED  
**Page 12**
- Moving the Manipulator and Polarizer Motors  
**Page 13**
- Using the Helium Lamp ...  
**Page 15**
- Setting the Undulator Gap and  
monochromator Wavelength...  
**Page 20**

## Overview of the Beamline:

Beamline 12.0.1.1 is the ARPES (Angle Resolved Photoemission Spectroscopy) branch of beamline 12 at the ALS. Below is a simplified diagram of the optical design of the beamline (See the end of this document for an enlarged version of this diagram). Light leaves the U8 undulator, after which it is collimated by the 4-jaw aperture before moving on to the first horizontal plane mirror (M1). It is then deflected slightly down by a spherical mirror (M2) to the monochromator. The monochromator contains two plane VLS gratings- one has 200 lines per mm, and the other has 1200 lines per mm. The 200 line grating is optimized for  $\sim 60$  eV and provides the resolving power of  $\sim 1200$ . The 1200 line grating is optimized for  $\sim 150$  eV with a theoretical resolving power of  $\sim 8000$ . The overall range of energies for this beamline is from about 20-320 eV. The beamline is quite bright- the 200 l/mm grating can deliver about  $10^{13}$  photons/s at 134 eV. This brightness made it possible to install the low energy circular polarizer which uses reflection off of four plane polarizing mirrors to turn the linearly polarized light from the undulator into left or right hand circularly polarized light. With this polarizer it is possible to achieve nearly 100% circularly polarized light for energies up to 80 eV. Just downstream from the monochromator grating is the exit slit, which can close down with precision to a few microns. Down from the exit slit is the M3 horizontal mirror, which deflects the beam to any of the three endstations at beamline 12, and controls the horizontal focus of the beam in the main chamber. Just downstream from the M3 mirror is where the circular polarizer is located- when not in use, it allows the beam to simply pass through to the vertical focusing mirror. Tweaking the vertical and horizontal mirrors should rarely be necessary- the beam has been aligned to maximize flux and resolution to all three endstations on Beamline 12.

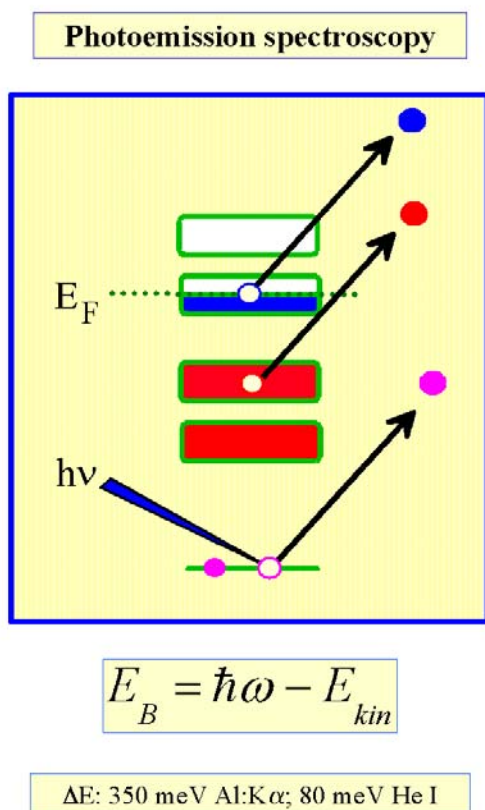


**Optical Design of Beamline 12.0.1.1**

## Photoemission Spectroscopy:

Photoemission Spectroscopy relies is a combination of two techniques of great importance in physics. The first is photoemission, whereby one may strike a surface with a photon and expel an electron with a characteristic energy, and the second is spectroscopy, where one studies the distribution of energies emitted from a sample, surface, or object. An excellent description of how the two work together is given by Laurent Alvarez on his personal website ([http://www.geocities.com/ CollegePark/3972/index.html](http://www.geocities.com/CollegePark/3972/index.html)):

“Photoemission is widely used in the study of the electronic structure of solids. It utilizes the photoelectric effect in which an electron is ejected from the occupied electronic levels of the sample. In a photoemission experiment, the kinetic energy of the photoelectrons usually varies from a few electron volts up to a few hundred electron volts, depending on the photon energy used. This results in the surface sensitivity of the technique, as the inelastic mean free path of a typical photoelectron in the solid is in the range of 5-30 Å. This means that UHV (ultra-high vacuum) is necessary to maintain a surface of adsorbates during the time scale of measurement, and that the surface effect should be borne in mind during the interpretation of the resulting spectra.”



The figure to the left shows “excitation of an electron from an electronic state below the Fermi level by X or UV rays (occupied states)”

Using the formula to the left, we may analyze the kinetic energy of the emitted electrons to get the binding energy of the state, telling us where the electron was excited from, or what materials may be present in the sample. The resulting spectra will give us insight into the density of the electronic state(s) that are present in the sample.

A more accurate and representative idea of what is going on with electrons within a solid and the angle-resolved aspect of ARPES is well described by the UK Surface Analysis Forum web site ([www.uksaf.org/tech/arups.html](http://www.uksaf.org/tech/arups.html)):

“In an isolated atom, the electrons occupy certain energy levels. In a solid, the levels involved in bonding ‘the valence states’ become blurred, and electrons can occupy a

range of energies bands. The energy of an electron in the solid depends on its momentum. Hence, by detecting photoelectrons emitted from a surface at different emission angles, the energy of the electrons as a function of the momentum vector may be determined. This process is

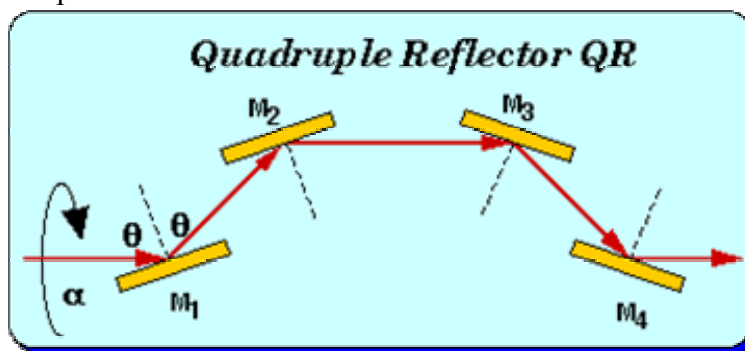


known as ‘band mapping’ and is a powerful probe of the electronic structure of crystalline materials. The measurements can usually be compared with theoretical predictions.”

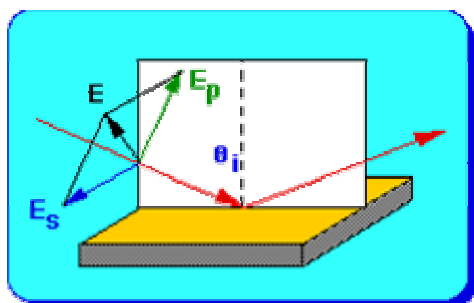
For information on how the SES-100 Analyzer works, please refer to its manual, located at the beamline.

### The Circular Polarizer:

The Circular Polarizer (or CP) at BL 12 was installed in September of 2002, and has opened up a broad range of experimental possibilities at the beamline. The CP can transform the linearly polarized light produced by the U12 undulator into elliptically polarized light of any orientation. More specifically and importantly, it can transform the light into left or right-hand circularly polarized light. It does this using the simple concept of plane reflection in a precisely aligned system of 4 mirrors.



**Optical Design of the Circular Polarizer**



**Reflection of an EM wave off of a metallic surface**

When light reflects off of a metallic surface, it is reflected into two resulting components, called the s- and p-polarized components. This is true for incoming light of any polarization. Reflection off of a metallic surface also introduces a partial phase change between the periods of the s- and p-polarized components. For each successive reflection off of a mirror, the phase difference between the s- and p- components changes.

This phase difference changes as a function of the angle of incidence of the light onto the reflecting surface (mirror). Since the CP has been designed so the beam reflects off of each mirror at the exact same angle, we produce 4 consecutive equal phase shifts. If we align our angle of incidence such that the total phase shift introduced over the four reflections is 90 degrees, we have perpendicular components of light with equal maximum amplitudes and a phase difference of 90 degrees- this is the definition of circularly polarized light.

#### Criteria for circularly polarized Light :

a) match Amplitudes

$$\tan \alpha = R_p^* / R_s^*$$

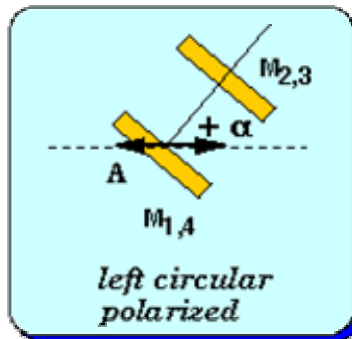
b) match total Phase Shift

$$\Delta = 4\delta = \pi / 2$$

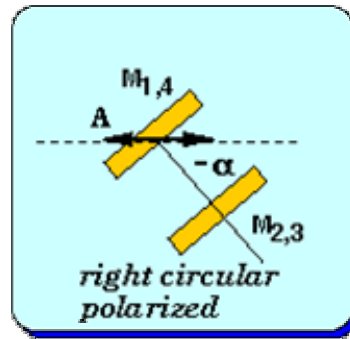
Since the CP is designed to send light out on the same axis as linearly polarized light into circularly polarized light by adjusting the angle of incidence, we can introduce a 90 degree phase difference. We adjust the alpha angle of the polarizer to choose either left or right circular polarization. For any given alpha angle, one can change theta in order to get CP light. Depending on whether the phase difference is + or – 90 degrees, we will have

#### **CP Light Criteria**

light that is either left or right circularly polarized- rotating alpha once we find either left or right CP will allow us to switch to the other.



**Left CP  
Mirror  
Alignment**

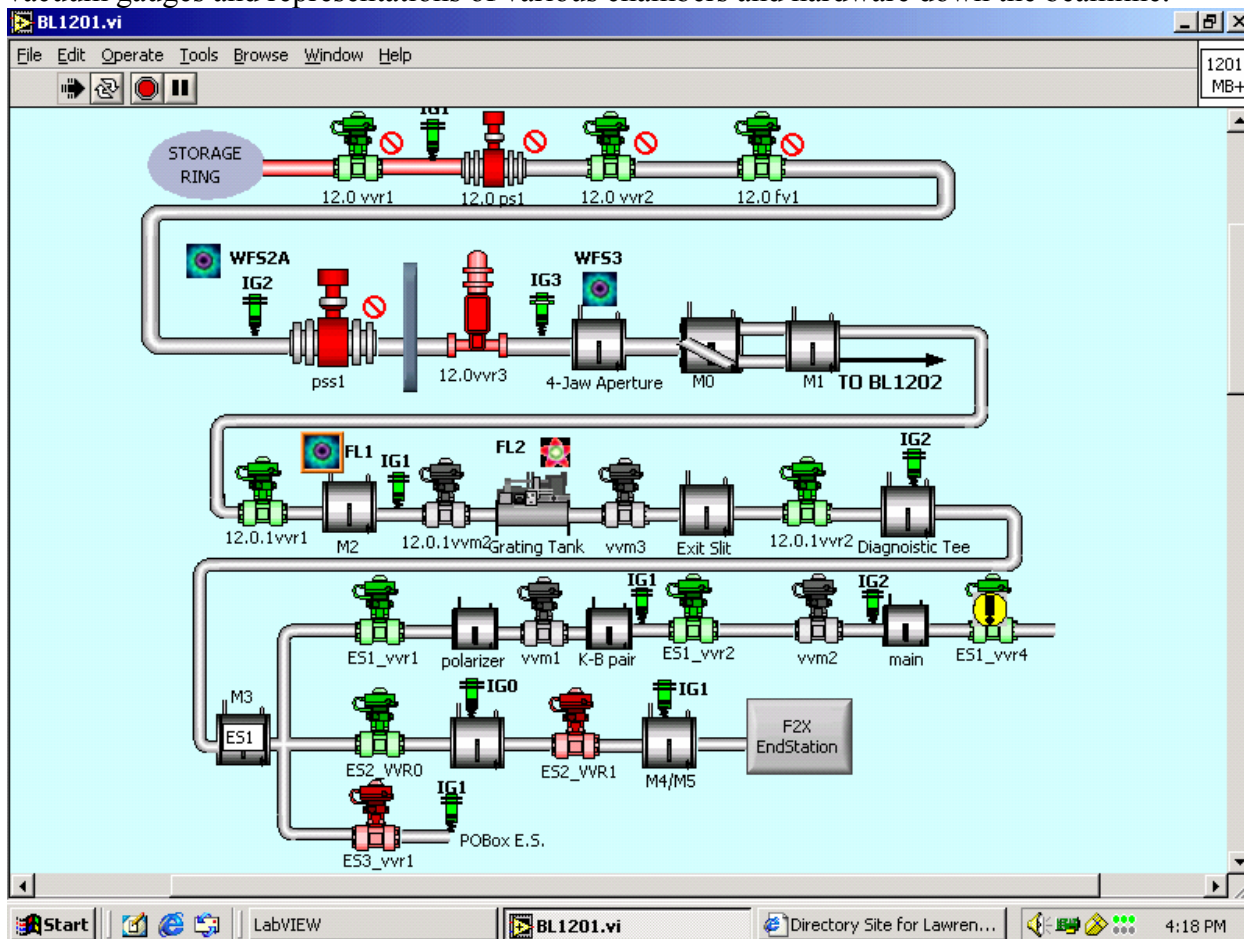


**Right CP  
Mirror  
Alignment**

A polarizer which works like this is referred to as a  $\lambda/4$  phase shifter or retarder. An excellent reference site for this circular polarizer is the page of Hartmut Höchst from University of Wisconsin-Madison, where the CP was built (<http://hhochst.src.wisc.edu/HHH.html>).

## Vacuum Safety Interlock System

This beamline (like all beamlines at the ALS) has a set of interlock systems to ensure the safety of users as well as prevent damage to the beamline and storage ring in case of accidents. The main safety interlocks are controlled by a LabView VI shown on the screen in the rack labeled BL1223 next to beamline 12.0.1.2. This program allows you to control all of the valves upstream from the endstation, as well as most of the valves on all of the three BL 12 endstations. The panel will show you a schematic drawing all of these vacuum valves, plus the various vacuum gauges and representations of various chambers and hardware down the beamline.



**The LabView VI to control the valves at Beamline 12**

During operation, open valves are colored green, while closed valves are colored red. All of the gauges, labeled IG1, IG2, etc. should be green. The schematic will also indicate how far the beam is passing through the tube by showing sections of the pipe which are open to the beam as red. Gray pipe indicates empty pipe. (Note: Pipe coloring on the schematic is solely a function of which gauges are opened, the schematic does not actually know if beam is passing through tubing). In order to get beam to the 12.0.1.1 endstation, for safety first make sure all valves downstream from and including the one labeled 12.0.1vvr2 are closed. If the beam is going to BL 12.0.2, first click of the figure in the schematic labeled M0, and switch the mirror to setting

12.0.1. Next, click on the figure in the schematic labeled M3, and switch the mirror setting to M1. The schematic will display three periods (...) while the mirror moves into place. Once the label M1 comes up on the figure, the mirror is in place, and you may open up the valves from 12.0.1vvr2 down. When beam is not available, you will not be able to open the main valve. If light is available to users, and the valves do not open, there is a problem or a lockout by EH&S, in which case you should call Alexei Fedorov (x7521), Rudy Kimmerling (x7519), or the control room (x4969). There is one upstream shutter and 2 manual valves NOT controlled by the VI. The shutter is located just upstream of M3, and is controlled by a small switch in the beamline equipment control rack labeled BL1225.



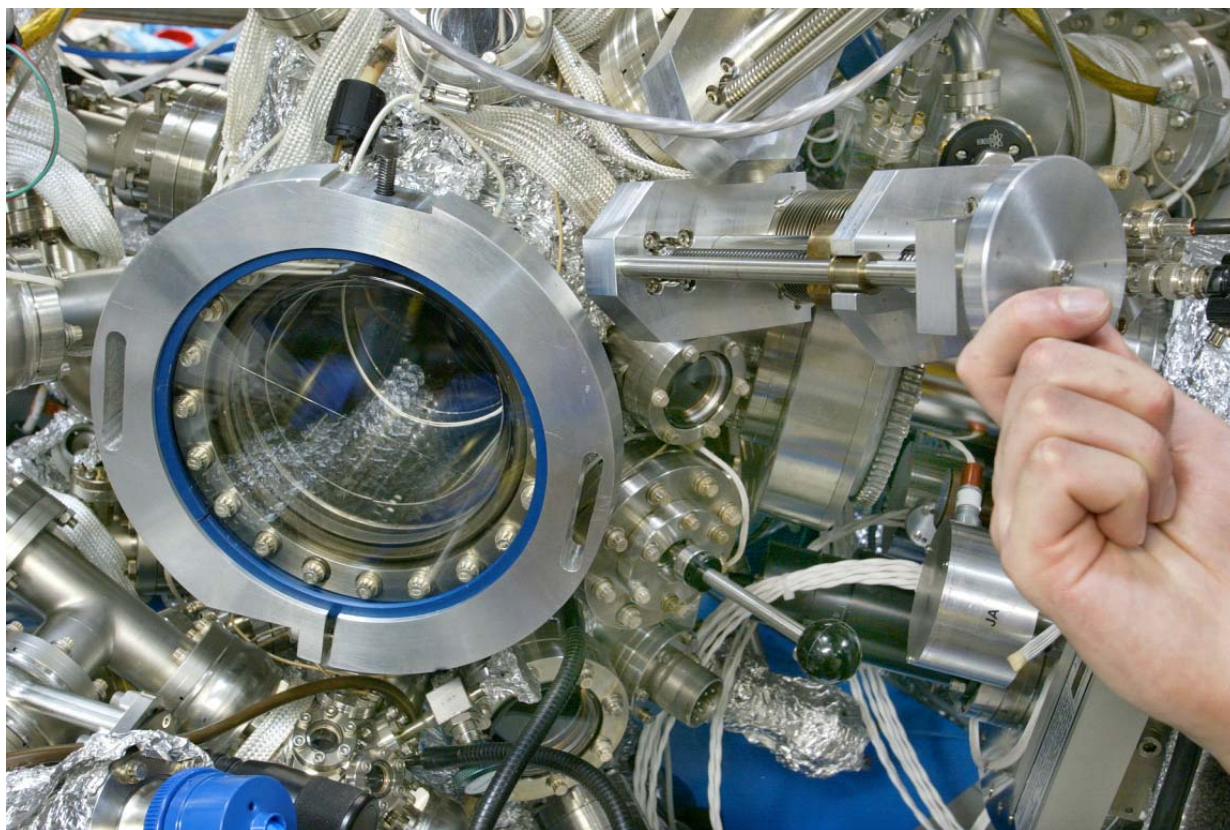
**Shutter Switch at BL 12 (left of green BNC cable, below round knob)**

It is labeled, and is open when the switch is in the down position. The first manual valve is just downstream of the circular polarizer, while the second one is just upstream from the main chamber, under the He lamp (these valves are usually left open). If all valves are open, and the VI shows red pipe all the way to the end, then beam should be available in the main chamber.

### **Beam Detection**

Beam should be easily detectable in two different ways. To check for beam, first raise the manipulator out of the way using the “Withdraw Manipulator” setting in the motor control software (discussed below), removing the sample from the path of the beam. A phosphor-coated window at the end of the chamber should show the presence of beam. Also, a YAG crystal that will illuminate can be moved into the beam path by rotating the handle located just to the right of the main viewport and above the handle used for cleaving the sample (see picture, next page). If you insert the handle with the YAG crystal further into the chamber, a photodiode will move into the beam path, which can be used to indirectly measure the photon flux (the resulting current will be directly proportional to the photon flux).





**The YAG crystal/photodiode insertion device**

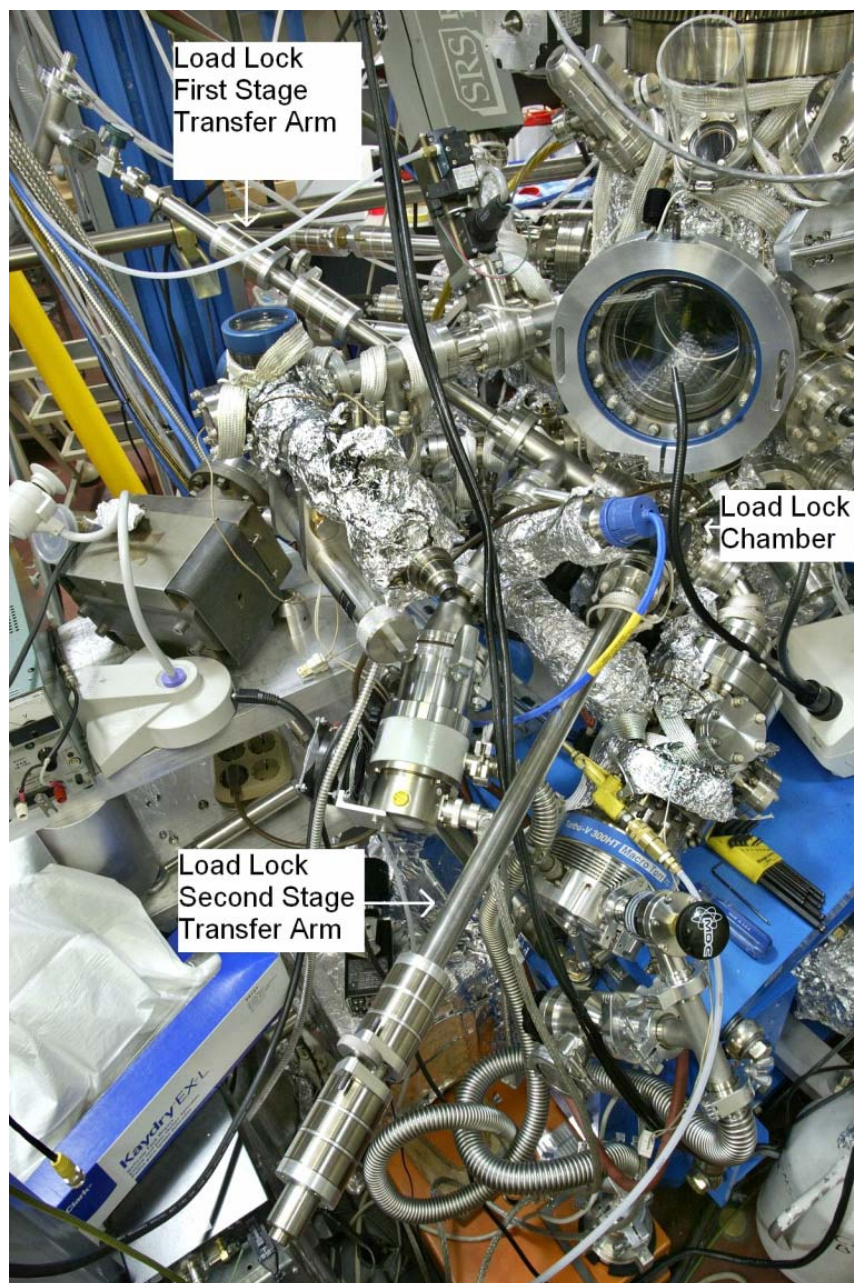
### **Swapping the Sample**

Inserting a sample into the chamber is a fairly straightforward, but delicate task. There are two stages of load locks in order to prevent excess gas from entering the chamber. Always make sure your sample is UHV clean before you put it in the chamber, as dirty samples may contaminate the chamber and cause delays.

To remove the sample

1. Use the motor control program to move the manipulator to “Sample Transfer” position in the one-touch trajectory menu.
2. Use the small flashlight attached (or another light source) to the small load lock chamber to look inside. This is the load lock second stage chamber. There is a small sample holder with two slots. Below the chamber is a two part rotating handle- the bottom part rotates the sample, and the top part of the handle raises and lowers the sample holder. Lower this holder out of the way of the transfer arm of the sample grabber.





**The 2-stage Load Lock system at Beamline 12.0.1.1 which allows for transfer of a sample from atmosphere to UHV pressure**

At this point, you should also make note of the pressure of the various stages of the Load Lock. The gauges are on the rack to the left of the prep table, with most of the Varian turbo pump controllers. The Load Lock 2<sup>nd</sup> stage pressure should be in the  $10^{-9}$  or  $10^{-10}$  range, while the 1<sup>st</sup> stage Load lock should be in the low  $10^{-7}$  or  $10^{-8}$  range.

3. Make sure that the small load lock 2nd stage chamber is valved off from the first stage load lock by closing the valve on the left side of the 2nd stage if it is open, and then open the valve between the main chamber and 2nd stage load lock chamber.
4. Using the viewport just to the right of the load lock chamber to look inside, insert the grabber and carefully remove the sample from the cryostat. Withdraw the grabber with the sample all the way back, and close the valve to the main chamber.
5. Raise the two-slot sample holder back up, and using the grabber, insert the sample into one of the two slots.

If the next sample is ready in the other slot of the holder, simply repeat this process in reverse to put the new sample in, remembering to valve off the load lock from the chamber after the sample has been inserted into the cryostat.

To remove the sample from the endstation completely:

1. After sliding the sample into the two-slot holder, first rotate the holder 90 degrees clockwise. Making sure that the load lock stage 2 is valved off from the main chamber, you can now transfer the sample to load lock stage one.
2. Open the valve between the 2 stages, which is located just to the left of the 2nd stage load lock chamber. Extend the first stage sample grabber down, grab the sample, and withdraw it all the way up and back into the first stage of the load lock.
3. Close the valve between the two stages of the load lock, and close the valve between the turbo pump and the first stage load lock. You are now ready to vent the first stage of the load lock.
4. Attach the nitrogen tube to the back end of the first stage load lock with a quick flange, turn on the nitrogen, and vent the load lock stage. There is a green Swagelok valve on the arm, make sure it is open while the first stage is being vented. When the stage is vented, nitrogen will flow from the release valve, and you can unhook the nitrogen tube.

You can now simply detach this part of the load lock at the flange above the 1st stage turbo. Be careful to withdraw the grabber straight out so you do not bang anything.

## Inserting a new sample

1. Once you have placed your new sample in the grabber, carefully reattach the transfer arm to where you removed it. A rubber gasket with 3 nuts and bolts is sufficient.
2. Once you are attached and tightened, attach the scroll pump located at the base of the endstation to the 1st stage load lock with a quick flange. Make sure the green handle valve is open, and turn on the scroll pump.

You will have to wait about 15 minutes for the scroll pump to get the 1st stage load lock down to a vacuum of about  $10^{-4}$ .

1. After about 15-20 minutes, close off the green valve, and shut down the scroll pump.
2. Close the valve on the back of the turbo pump for stage 2 (to prevent a backflow of air into the second stage pump, slowing it down and lifting the pressure), and open the valve between the turbo pump and first stage load lock.

3. After a minute or so, when the 1<sup>st</sup> stage pressure reaches the mid to high  $10^{-6}$  range, open the valve on the back of the turbo pump for phase two again.

Once the pressure gets down to the mid  $10^{-8}$  range, it is safe to transfer the sample from the first to the second stage, using the reverse process as described above.

### **Venting the Main Chamber**

The main chamber is best kept under high vacuum at all times; however, sometimes it is necessary to vent it and open it so that items may be placed inside or on the chamber. Always contact Alexei Fedorov (x7521) or Rudy Kimmerling (x7519) before venting the chamber, as doing so will cause a delay of at least two days for bake-out before the main chamber can be opened back up to the beam. In the event that you must vent the chamber, the first step is to make sure you close every valve surrounding the chamber. First, use the computer program on the panel (see Vacuum Safety Interlock System) to close all valves surrounding the main chamber. To minimize the amount of apparatus that needs to be baked, and to avoid damaging the high-vacuum pumps, you must also valve off all of the pumps and hand valves connected to the chamber. Make sure to close the following valves: Directly upstream from the chamber, there is a hand controlled valve, it is directly under the Helium lamp, and a little hard to reach. Also, make sure that the hand valve between the chamber and load lock is closed. Close the hand valve directly beneath the He lamp, if it is not already shut. Most valves that isolate pumps are controlled by buttons located on the top shelf of the portable blue rack of controllers to the left of the chamber (see picture, next page). Make sure all four valves (main turbo, ion pump, cryo pump, He monochromator) are closed. There are three more button controlled valves on the box in the rack above the workbench numbered BL 1231. Make sure that these three valves, VVR2 Main/KB, VVR4 LL/Prep and VVR4A Vent LL, are closed. All valves should now be sealed, and the main chamber should be isolated from the rest of the apparatus. Near the foot of the blue rack is one last gray box that controls the valve for the Gd pump to its left, close it as well. Approximately 1 meter below the Scienta analyzer is another large hand operated valve which isolates the second turbo located underneath the chamber, turn this handle clockwise all



the way until it is in the closed position. Just to the left and above this pump is the vent that is attached to the main chamber.

**The Valve to Vent the Main Chamber**



If the scroll pump is attached, make sure that the valve just to the right of the vent is all the way closed. Remove the scroll pump and attach the nitrogen hose to the vent with a quick flange. Start the flow of nitrogen using the regulator located above the KB chamber of BL 12.0.1.2. There is a relief valve on the vent; when you feel the nitrogen flowing, turn off the ion gauge controller for the main chamber, and open the valve just to the right of the vent. You may need a wrench to do so. The flow of air from the relief valve should stop for a few minutes; when you feel it once again, the chamber is vented, and is safe to open.

## **Using the LEED**

In order to obtain a good LEED image, first you must move the sample into place. Move the manipulator into place by using the motor control software and the one-touch trajectory called "LEED." The sample should now be in place. The camera is connected to the same cords as the analyzer camera, simply remove the wires and attach to the LEED if it is not already done. Switch on the monitor in the blue rack, switching to channel A to see the camera image. The controls for the LEED voltage, the Omicron Vakuumphysik GMBH, are in the same rack, below the two Ionization gauge controllers. Turn it on, and set the following reference voltages:

With a reference Energy

$E=333 \text{ eV}$

$I \text{ filament} = 1.78 \text{ A}$

$I \text{ emission} = 2.436 \text{ mA}$

$W = 108$

$L \text{ } 1/3 = 213$

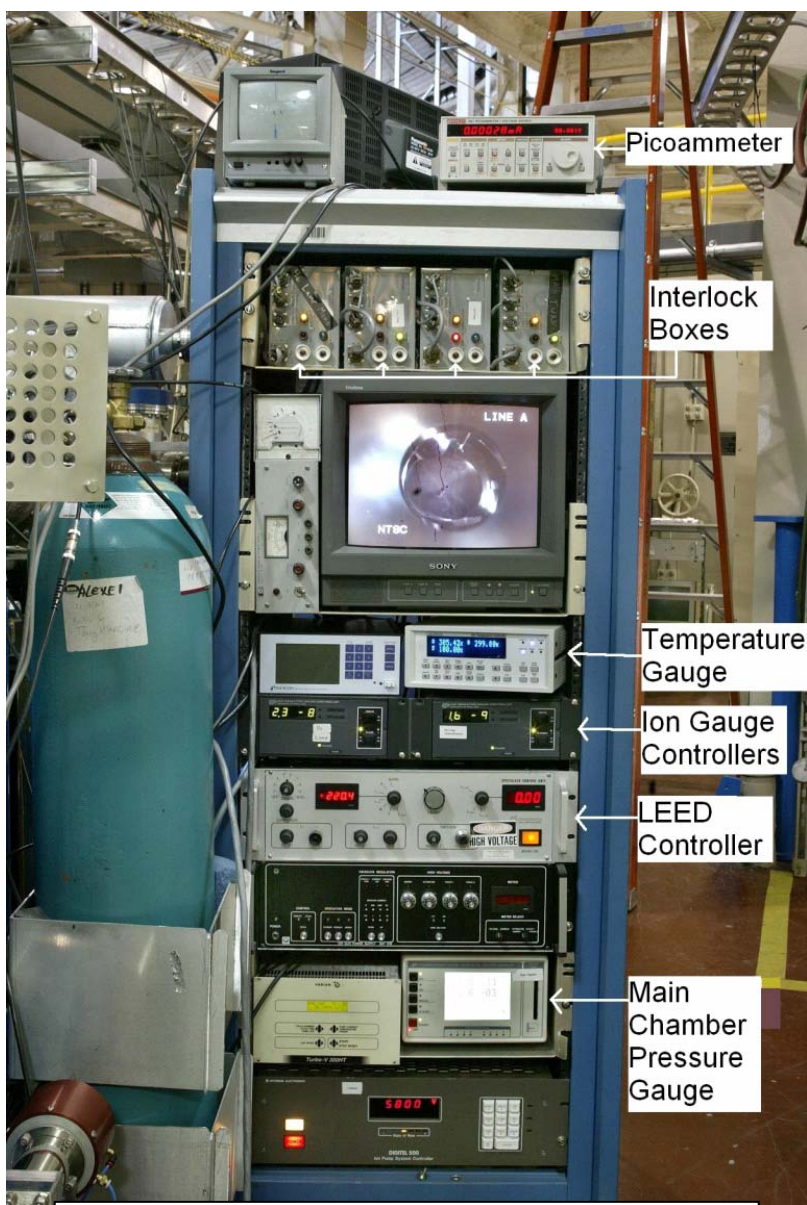
$L2 = 50$

$A = 329$

$\text{Suppr.} = 44$

$\text{Screen} = 10 \text{ kV}$

(Note: With these voltages, the LEED gets really bad below 290 eV)



**The Blue Rack and its Controllers/Gauges**

### **Settings for Energies below 290 eV**

LEED for E= 170.4eV

L 1/3= 131

L2= 34

A= 328

At E= 60 eV, LEED is very bad, but the best voltages are at:

L 1/3= 128

L2= 35

W= 108

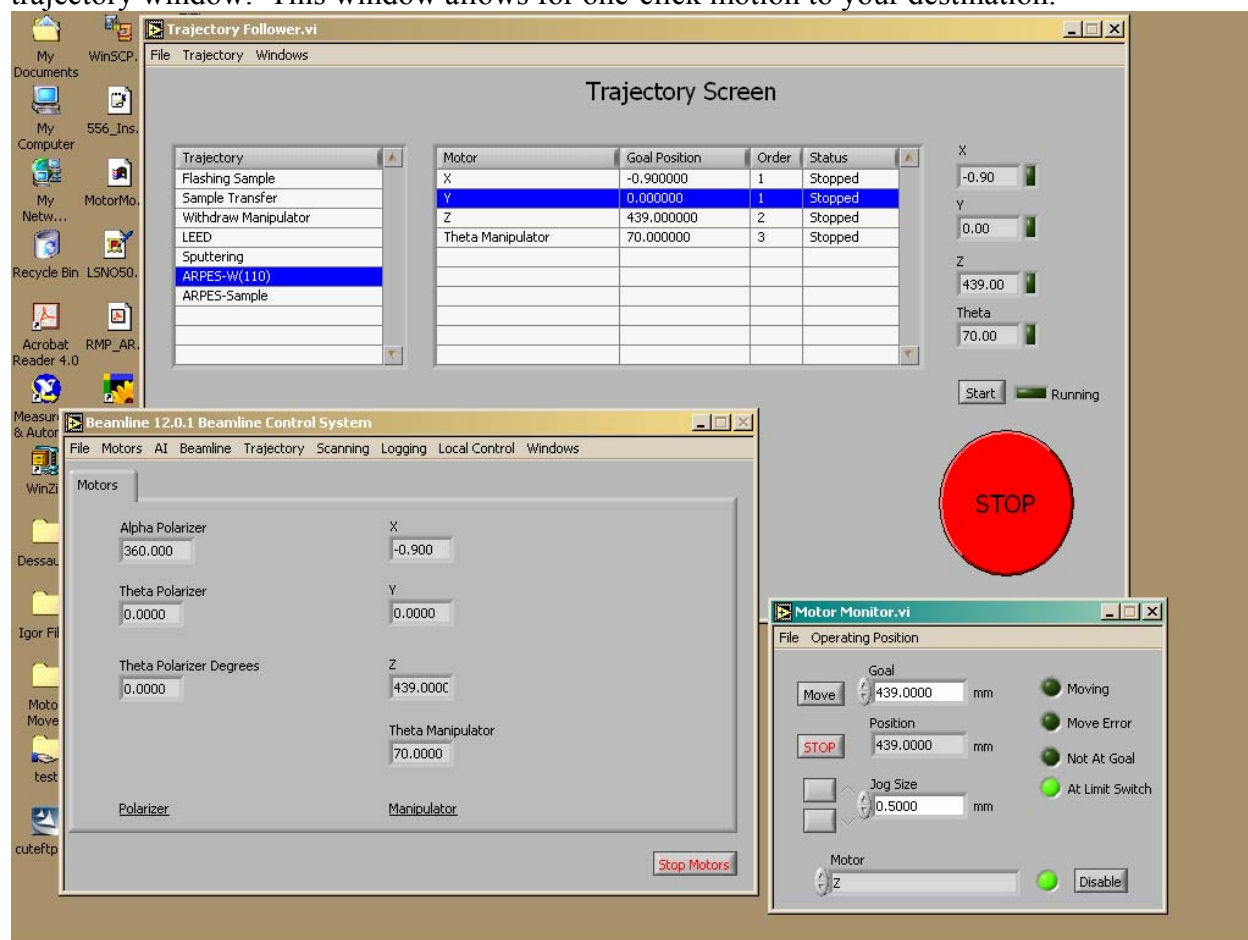
A= 330

In order to rotate the sample post itself about its axis (to rotate the image on the screen around the center of the screen) there is a dial attached to a screwdriver that is inside the chamber. This dial is a bit hard to reach; it is located just to the left of load lock chamber 1. Rotating the base brings the screwdriver closer or further towards the sample back, and rotating the dial itself spins it. Slowly and carefully move the screwdriver into place, rotate the dial, and be careful not to push the entire sample forward. When you have reached the optimal position, make sure to withdraw the screwdriver all the way back out of the way from the sample, so it does not strike the cryostat as it is raised, lowered, or rotated.

### **Controlling the Manipulator and Polarizer Motors**

There are 4 degrees of freedom in the motion of the manipulator, X, Y, Z, and Theta. There are two simple ways to move these motors, and the manipulator. One way is to use the joystick, a gray rectangular box with 6 red buttons, 2 black buttons, and a switch. The switch controls the jog velocity. The top pair of buttons, labeled motor 1, controls the theta motor. The pair labeled Motor 2 control the Z motor, the lowest red pair controls the Y motor, and the black pair controls the X motor. Using the joystick, however, ruins the backlash compensation and is not precise in terms of its motion. **Do not use the joystick unless the software to move the manipulator is broken.** The second way to control the motor, and to monitor its position, is using the software on the white Vectra computer at the beamline. There is a shortcut to the LabView VI that performs this operation on the desktop labeled "BL Control Main." When you open this program, you will see two columns of indicators, one for the polarizer, and one for the manipulator. To run the VI, press the little white arrow in the top left corner. A second window will pop up with a small box to send the motor to a given position or to jog it. When you do this, a smaller window will also open displaying three indicators and a few buttons and controls. This window, labeled "Motor Monitor.vi" has a small control on the bottom with each motor's name in it. You can scroll through to control any of the 6 motors at the beamline. This VI is fairly self explanatory; you simply enter your requested position, and press the Move button. No home position is specified, but soft limits should be in place to ensure that the manipulator is not damaged. Still, be careful when jogging or setting target distances, and do not continuously

press the jog button, this tends to jam the program. The newest addition to this program is the trajectory window. This window allows for one-click motion to your destination.



### The Beamline 12.0.1.1 “Beamline Control System” Software

Several standard positions, such as sample transfer, LEED, etc. are programmed into this window, and you can get to virtually any important position at the click of a button. Please do not edit the trajectories yourself- if you find the positions to be off, use the jog buttons in the program to move to the correct spot, and make a note of the new position for the beamline scientists (Alexei Fedorov and/or Rudy Kimmerling). The program should be run continuously so you can monitor the position- but position will stay recorded if you shut down the program. The program only needs to be reset if the controllers themselves lose power. The angles represented on the program are calculated from number of steps sent- there are no encoders- so it is always best to physically observe where the manipulator is if you can. In order to stop this VI, which should not be necessary unless the program seems to be malfunctioning, simply go to File-> Stop Beamline Controls. If you would like to change the parameters of the motor motion, please speak with Alexei or Rudy first, such elements as speed, acceleration, backlash, and gear ratio have been optimized given the needs and limitations of the system.

On the polarizer, there are 2 parameters you can control. The first is the tilt of the 4 mirrors, which is represented in the polarizer controls as theta, or theta degrees. The second is the

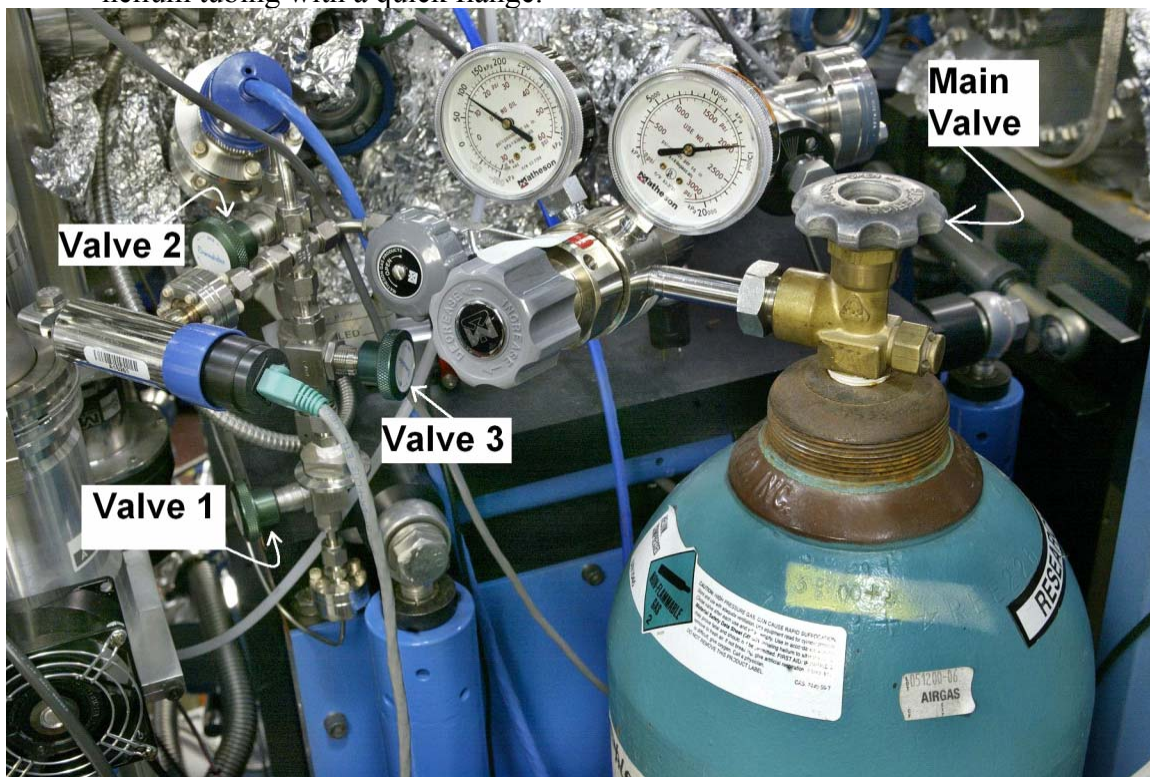


rotational angle of the mirrors about the beam's axis. This parameter is designated as Alpha. Press on the control in the Motor Monitor window and scroll to either Theta Polarizer Polynomial to control theta, or Alpha Polarizer to control alpha. DO NOT USE THE "THETA POLARIZER" SETTING UNLESS YOU WISH TO JOG THE MOTOR A FEW STEPS. When you have chosen your motor, simply type in your destination value in the top "Goal" control, and click the move button. The position indicator will show you the current position of the motor. Although there are many features to this VI, you should not need to go any further than the Motor Mover window. To stop the VI, go back to the first window called Beamline 12.0.1 Beamline Control System, and click on File-> Stop Beamline Controls, which should automatically close the Motor Mover window and stop the VI.

### **Using the Helium Lamp**

The Helium Lamp on BL 12.0.1.1 allows for data taking even when ALS beam time is not available. There are many steps to follow in order to ensure proper usage of the He lamp, as it is a delicate instrument. The first step is to remove the old helium from the system and flush it with fresh helium before use.

- First, ensure that the valve on top of the green helium tank is tightly closed. Attach the scroll pump which is at the base of the endstation to the vent at the helium tubing with a quick flange.



**The Helium Tank, regulator, and valves for the BL 12 He lamp**

few seconds for the He to pump out.

- Close valve 3, and open the main valve atop the green tank, putting fresh Helium into the system.
- After a few seconds, close the main valve on top of the green tank, and open green valve number 3 again. Let it pump for a few seconds, then close green valve 3 again tightly, and turn off and remove the scroll pump from the vent.
- Open the valve on top of the green tank, and adjust the valves until the pressure in the left side of the regulator is at about 10-13 psi. You now have fresh He in the system.

The next stage of the process involves getting the right He pressure inside the lamp.

There is a gauge to measure the He pressure which sits in the blue rack and is labeled “He Lamp.” Normally, this pressure should be in the  $10^{-8}$  range. Check the pressure, and make sure that the two hand valves attached by the lamp are closed- one separates the small turbo pump from the lamp, the other separates the lamp from the monochromator.



Open the small leak valve slowly while watching the pressure, and adjust it to the point that the pressure goes to about  $2$  to  $3 \times 10^{-4}$  Torr. Make sure that the pressure is not above  $6 \times 10^{-4}$  Torr, or else the lamp could be damaged when the microwave generator is on. If the pressure is below  $2 \times 10^{-4}$ , it will be harder to start the lamp.

**The He Lamp and He Leak Valve**



It is now time to turn on the lamp with the Gammadata microwave generator located in the rack labeled BL1124, which is directly across from the chamber.

- First, walk around and make sure that the door behind the Gammadata is open; the Gammadata has a fan which vents out the back. You should not run the Gammadata with the door closed. When it is open, come back to the front and switch on the orange power button.
- Next, press the black UHV ON button. The two green lights, Filament On and Filament OK will go on, which means everything is normal, and the microwave generator is starting up.

In about 2 minutes, the voltage meter on the Gammadata will bounce up, as will the discharge current and reflected power. The discharge current should be between 250 and 275 mA, the voltage should be about 4.5 kV, and the reflected power should be below 10%.



**The Gammadata Microwave Generator**



- You want to minimize the reflected power, so if it is not zero, adjust the pressure once again with the leak valve, watching the reflected power meter to get the lowest possible reflected power. Check to see that everything is within these given okay ranges.
- When they are, first close the valve to the Cryo pump- He exposure may damage the Cryo pump.
- When you are ready, open the two hand valves, the one to the pump, and the one to the monochromator.

When you open these valves, you should also make sure that the valve for the larger turbo which pumps the monochromator is open. This valve is the pneumatic valve labeled "He Lamp monochromator" in the top left corner of the portable blue rack located right next to the large green bottle of Helium. Press the right (green) button to open it, it is okay if the lights do not work, the valve is still open.

- Upon opening the valves, you may once again have to play with the He pressure to minimize reflected power and optimize the settings. When you have done this, make a quick check of all of the pressures. The pirani gauge that is read by the second slot of the Stanford Research Systems gauge in the blue rack (the top slot of the same gauge is used for the main chamber pressure) is attached to the roughing pump that gets fed by all three turbos on the He lamp system. When there is no He present, it should read below 5 mTorr pressure, but with Helium in the grating tank and the valves open, it should read between 60 and 100 mTorr pressure.

Now it is time to tune the monochromator to get the correct line of Helium. The standard setting of the monochromator should have the He 1 line in focus. However, if the He 1 line does not appear, then it is necessary to play with the monochromator.

- First off, check the pressure in the monochromator chamber on the gauge below the Sony monitor on the right side of the blue rack labeled "He Lamp monochromator." The pressure should be about  $3 \text{ or } 4 \times 10^{-6}$ . We need to know which line of Helium we are looking at, so we must move the photodiode into the path of the beam. The photodiode is located on the same rod as the YAG crystal described in the section "Beam Detection."
- Move the photodiode into place using the process described in the Beam Detection section, simply rotating the panel until the YAG crystal moves past the beam, and the photodiode moves into the line of the beam. The picoammeter located on top of the blue rack should display the current in micro- or nanoamperes. Make sure the picoammeter is hooked up to the photodiode by connecting it to the left most BNC connector next to the handle for inserting the YAG crystal/photodiode. One of the other BNC connectors should be grounded to itself.

- As you move the photodiode into the path of the beam, watch the picoammeter and stop rotating where the current is maximized. The monochromator should be set on the He 1 line, and should register about 2-3 microamps on the photodiode.
- If it does not, you adjust the monochromator using the screw handles on its back side. There are two handles. The one offset from the center of the flange controls the tilt and therefore wavelength of the monochromator.



- Play with this to find the line- if you are having trouble, move it all the way out (turn it counter clockwise/left) to find the zero order, which should be very bright. Then, move it back. The first line you should see is the He 2 line. The second is the He 1 satellite line, and finally, the third should be the He 1 line. Check to see that the current is between 2 and 3 microamps.

### Controlling the alignment of the He Lamp Monochromator

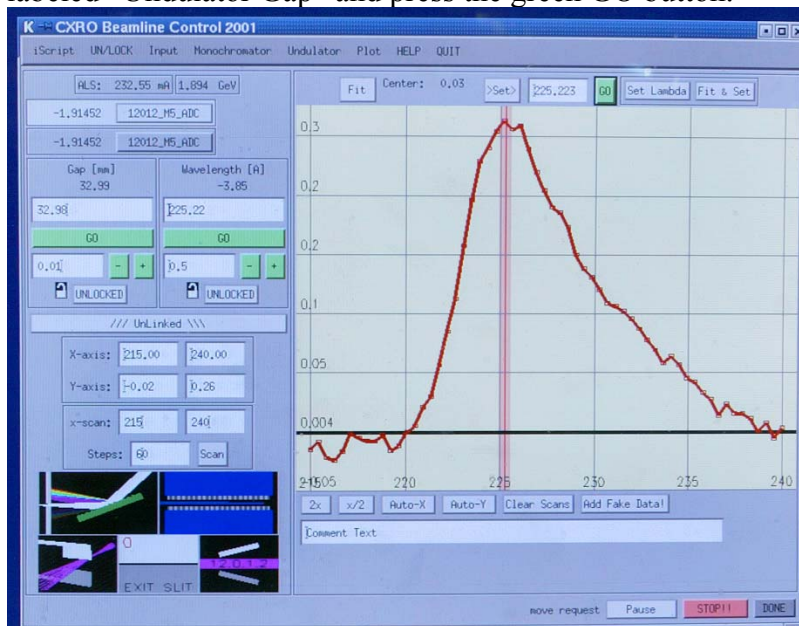
- The other handle, in the center of the monochromator, will steer the beam further left or right, so be careful. Remember that making adjustments to increase the flux or change the pressure will affect the reflected power, which should be kept as low as possible while still allowing a little play to maximize flux. Should the reflected power somehow exceed 10%, the microwave generator will shut down automatically. To restart it, simply check all of your connections and settings again to see that they are okay, and hold the UHV ON button down again until the green lights come back on.

\*At this point, a small note is to ensure that water is flowing to the He lamp to cool it. The two tubes leading to the lamp, one red and one green, supply cool water to and return it from the lamp. The small black valve should be open to allow water flow to the lamp. In general this is open, but it is worth taking a few seconds to check.

- **To turn the He monochromator off, simply press the UHV OFF button. Do NOT turn off the orange power button right away- the fan must continue to run for a while, or the Gammadata microwave generator could overheat and be damaged.**
- After you have pressed the UHV OFF button, close the He leak valve, close the two hand valves (but not the pneumatic He Lamp Mono. valve), and wait for the temperature to drop to  $\sim 25^\circ$  before turning of the main power with the orange button on the Gammadata.

## Setting the Undulator Gap and Monochromator Wavelength

The undulator gap and monochromator wavelength are set using the program on the computer by beamline 12.0.1.2, near the screen where valves and beam direction is controlled. On the left computer, you will see a program which allows you to control various hardware from the upstream part of the beamline. Type your desired gap width for the undulator into the small box labeled “Undulator Gap” and press the green GO button.

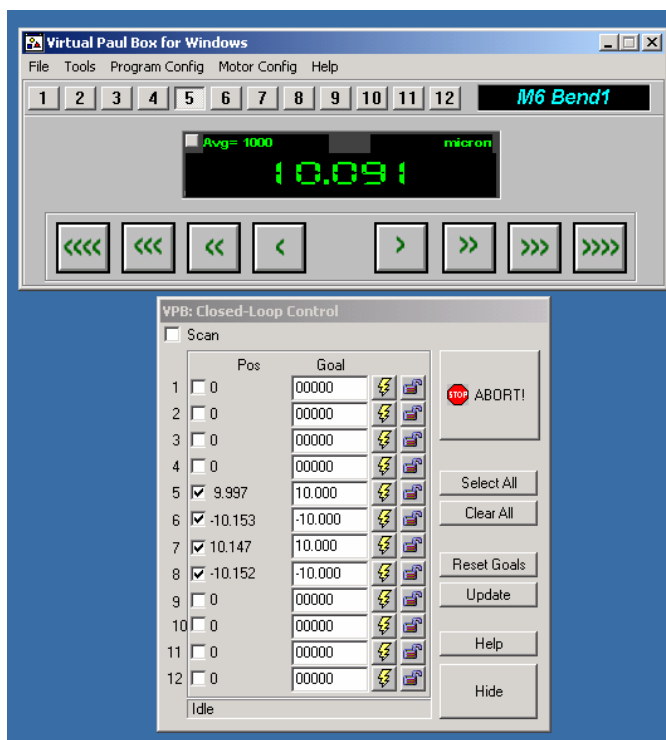


The gap width is shown in the program, and also can be seen throughout the ALS on the television screens displaying the Ring Status- in the bottom right corner, all gap widths are listed. This is a good way to double check the size of the undulator gap. The monochromator position, and therefore wavelength, may be adjusted in much the same way via its respective control and display in the program. It is also possible to take monochromator scans with this program.

## The Monochromator and Undulator Gap Software

On the computer to the right, you will see the controls for the exit slit width. The 4 positions of the respective corners of the slit are shown as positions 5, 6, 7, and 8. You can only change these one by one, so if you are going from a very small to very large gap, or vice versa, move incrementally (say first, move all motors from 170 to 50 microns, then all motors from 50 to 20 microns, and finally 20 to 5).

### The Software to Control Exit Slit Width





## Appendix 1: The Prep Chamber (Installed April 2003)

The prep chamber added to the load lock system in April 2003 is a convenient solution to the problem of high temperature annealing of samples in a low pressure environment.

Previously, there was no way to anneal a sample within the chamber without risking contamination of the chamber, or damage to the sample (due to the fact that it would have to be heated from the front). The new prep chamber allows for high temperature annealing outside of the main chamber, with simple two step load lock transfer to easily put the annealed sample into the chamber. The chamber is located between the 2<sup>nd</sup> stage of the load lock, and the 2<sup>nd</sup> stage turbo pump- just to the right of the load lock chamber. It had a single isolated slot for a special sample holder. Though the back end of the sample holder is the same, the front end consists of an elongated neck and horizontal plate to mount the sample on. These special sample holders are made of Molybdenum and have several special features. The top end crown of the holder has a wedge cut out for easier grabbing should the transfer arm be misaligned, and the horizontal plate has several holes for bolts- it is suggested that you mount your sample on a thin square of tantalum foil to be bolted onto the sample holder. Silver epoxy may contaminate the turbo pump if it is heated too fast- it is suggested that users use thin tantalum strips, spot welded, to hold their samples in place. When your sample is mounted, in order to load it simply follow the steps outlined above for sample transfer. Once you are ready to load the sample into load lock stage 2: Open the valve between phase 1 and phase 2.

Lower the two-slot sample holder far down, out of the way.

Lower the first stage transfer arm all the way down through stage 2 into the slot for the sample in the prep chamber.

Before you release the sample holder, rotate it until the sample itself is facing the window and fully visible, so the heater will align correctly with the back of the sample holder.

Withdraw the transfer arm all the way back to the first stage, and close both hand valves between the various stages. This will prevent contamination of the second stage of the load lock should anything go wrong.

## Appendix 2: The Helium Lamp

Slight modifications have been made to the He lamp over the past few months- quick valves were replaced by welded pieces and better swagelock pieces were fitted into the system. Due to this, the process for cleaning old helium out of the system is slightly different. It is now best to pump from the valve located up by the lamp itself, this way old helium is not pumped through the filter. Locate the valve, and pump the helium in the system out with the valve to the filter setup closed.